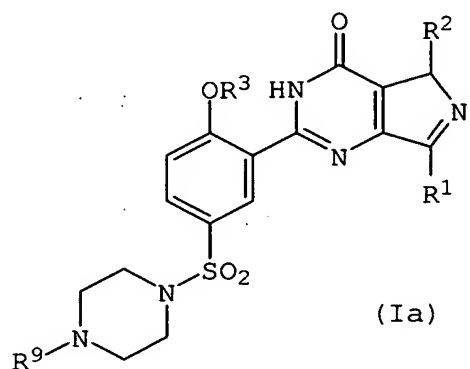


IN THE CLAIMS:

1. (Cancelled)

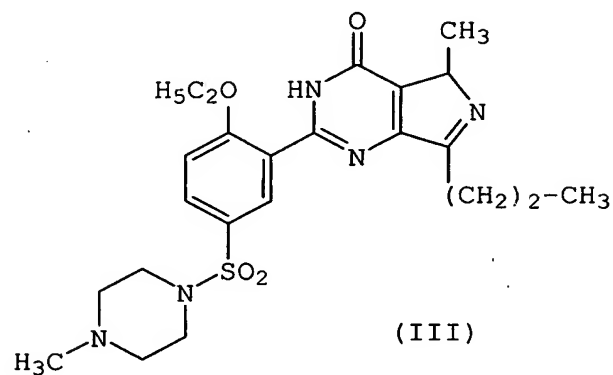
2. (Previously amended) The method of claim 5 wherein the pharmaceutical agent comprises a compound of formula (Ia):



wherein R<sup>9</sup> is an alkyl group having 1-4 C atoms which, optionally, are substituted with halogen or replaced by halogen;

or a pharmaceutically acceptable salt thereof.

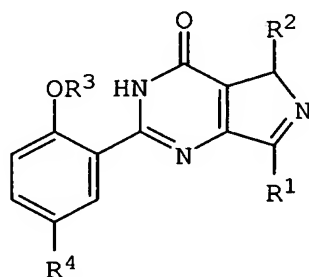
3. (Previously amended) The method of claim 5 wherein the pharmaceutical agent comprises a compound of formula (III):



or a pharmaceutically acceptable salt thereof.

4. (Cancelled)

5. (Currently amended) A method for a chemotherapeutic treatment of ~~neuropathies~~ an autono-  
mous neuropathy characterized by application to a  
 patient in need thereof of from 1-100 mg/day of a  
 pharmaceutical agent comprising a compound of formula  
 (I):



(I)

in which

$R^1$ =C<sub>1-6</sub>alkyl, optionally substituted with  
 halogen,

$R^2$ =hydrogen or C<sub>1-4</sub>alkyl, optionally sub-  
 stituted with halogen or replaced with halogen,

$R^3$ =C<sub>2-4</sub>alkyl, optionally substituted with  
 halogen,

$R^4$ =SO<sub>2</sub>NR<sup>5</sup>R<sup>6</sup>,

C<sub>1-4</sub>alkyl, optionally substituted with NR<sup>5</sup>R<sup>6</sup>,  
 CN, CONR<sup>5</sup>R<sup>6</sup>, CO<sub>2</sub>R<sup>7</sup>, or halogen,

C<sub>2-4</sub>-alkenyl, optionally substituted with  
 NR<sup>5</sup>R<sup>6</sup>, SONR<sup>5</sup>R<sup>6</sup>, CONR<sup>5</sup>R<sup>6</sup>, CO<sub>2</sub>R<sup>7</sup>, or halogen,

C<sub>2-4</sub>-alkanoyl, optionally substituted with  
 NR<sup>5</sup>R<sup>6</sup>, SONR<sup>5</sup>R<sup>6</sup>, CONR<sup>5</sup>R<sup>6</sup>, CO<sub>2</sub>R<sup>7</sup>, or halogen,

R<sup>5</sup> and R<sup>6</sup>, independent of one another, rep-  
 resent hydrogen or C<sub>1-4</sub>alkyl, or, together with the  
 nitrogen atom to which they are attached, represent a  
 pyrrolidino, piperidino, morpholino, 4-(NR<sup>8</sup>)-1-pipera-

zinyl or 1-imidazolyl ring which, optionally, may be substituted with one or two C<sub>1-4</sub>alkyl groups,

R<sup>7</sup>=hydrogen or C<sub>1-4</sub>alkyl, optionally, substituted with fluorine, and

R<sup>8</sup>=hydrogen, C<sub>1-3</sub>alkyl, or hydroxy alkyl having 1-4 C atoms, or a pharmaceutically acceptable salt thereof.

6. (Cancelled)

7. (Previously presented) The method of claim 5, wherein from 5-50 mg/day of said pharmaceutical agent is administered to a patient being treated.

8. (Previously presented) The method of claim 5, wherein from 25-50 mg/day of said pharmaceutical agent is administered to a patient being treated.

9. (Previously presented) The method of claim 5 wherein the neuropathy comprises a peripheral diabetic polyneuropathy.

10. (Previously presented) The method of claim 5 wherein the neuropathy comprises gastroparesis.

11. (New) The method of claim 5 wherein the neuropathy comprises a degenerative neuropathy.

12. (New) The method of claim 5 wherein the neuropathy comprises a toxic neuropathy.

13. (New) The method of claim 5 wherein the neuropathy comprises a metabolic neuropathy.

14. (New) The method of claim 5 wherein the neuropathy comprises an ischemic neuropathy.